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Experimental study of methane-oriented gasification of semi-anthracite and bituminous coals using oxygen and steam in the context of underground coal gasification (UCG): Effects of pressure, temperature, gasification reactant supply rates and coal rank

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Abstract:

The results of ex-situ small-scale laboratory tests performed in a bespoke batch reactor simulating coal gasification to find the most optimal experimental conditions for producing methane-rich syngas in the context of UCG are presented in this paper. The influence of gaseous reactants (oxygen and steam), their supply rates and thermodynamic conditions (temperatures of 650°C, 750°C, 850°C and pressures of 20 bar and 36 bar) on the gasification of semi-anthracite (South Wales coalfield) and bituminous (Silesian basin) coals is investigated. Increasing the gasification pressure from 20 bar to 36 bar and doubling the amount of steam with respect to oxygen benefit the methane generation. Although temperature increase from 650°C to 850°C also benefits methane generation, gasification at 750°C provides the most optimal conditions for methane-rich syngas production. Overall, the highest methane generation occurs at 750°C, 36 bar and H<sub>2</sub>O:O<sub>2</sub>=2:1 yielding peak methane concentrations of 44.00 vol.% and 35.55 vol.%, and average methane concentrations of 15.34 vol.% and 14.64 vol.% for the semi-anthracite and bituminous coals, respectively. These findings demonstrate that an increase in coal rank favours the methane generation. Owing to high methane content, the syngas produced at such conditions contains the highest calorific value, although the generation of hydrogen and carbon monoxide is reduced in comparison to the experiments conducted at 850°C. This study shows that gasification of bituminous and semi-anthracitic coals at elevated pressures can provide stable generation of methane-rich syngas whose quality can be controlled by the gasification temperature through the dynamics of steam and O<sub>2</sub> supply rates.

Keywords:

Underground coal gasification; Methane; Semi-anthracite; Bituminous coal; High pressure, High temperature.

## 1. Introduction

With an increasing global population, the society is faced with a dual challenge to produce more energy and less carbon. In 2018, the global energy consumption increased by 2.3%, i.e. at nearly twice the average rate of growth since 2010, while energy-related CO<sub>2</sub> emissions rose 1.7% [1]. Underground coal gasification (UCG) is a clean-coal technology that offers a prospect to be one of the solutions to such challenge, as it can safely harness the energy by utilising deep coal seams and convert them into a synthetic gas. Although coal reserves are the most abundant of all the fossil fuels, less than one sixth of the world's coal is economically accessible via conventional mining methods [2]. European coal reserves are mainly found in the UK, Germany, Poland, Netherlands, Ukraine and Turkey [3]. For instance, in the UK total coal resources could be 187 Btonnes with around 17 Btonnes suitable for UCG, representing a resource for almost 300 years [4, 5]. Similarly, Germany contains around 6.7 Btonnes of coal, while Poland has around 21 Btonnes of hard coal reserves alone [5]. As the global demand for gas is forecast to increase, with industry emerging as the main driver for such growth [6], utilisation of coal reserves through UCG offers a great prospect of increasing the efficiency of fossil fuel use and security of energy supply, and help moving towards low carbon emissions in Europe and worldwide.

The UCG process involves the reactions of oxygen, air or steam with coal carbon and other pyrolysis products producing a syngas which is mainly composed of methane, hydrogen, carbon monoxide, carbon dioxide and steam [7]. For example, hydrogen is a clean fuel that constitutes a valuable chemical feedstock for different chemical processes and syntheses [8]. The produced methane-rich syngas is compatible with natural gas and can be used as a synthetic natural gas (SNG) for power generation, raw material in industry or conversion into other fuels and chemicals [7, 9, 10]. In general, as a fuel, methane has a high economic value as well as the environmental, energy security and industrial safety benefits [11].

As coal is not a pure chemical species, but rather a heterogeneous and multispecies porous medium with large available internal surface area, its gasification involves a complex system of chemical reactions, with the main ones shown in Table 1 [7, 12-14]. In general, endothermic reactions are favoured at higher temperatures, at the expense of exothermic reactions [12]. Those reactions in which

the number of moles of products is less than the number of moles of reactants will be favoured at higher pressures [12]. Thermodynamically, some CH<sub>4</sub> producing reactions, such as hydrogasification reaction (no. 4) are favoured at high pressures and relatively lower temperatures [7].

Table 1. Main gasification reactions (after [13, 14])

No.	Reaction	Reaction enthalpy (MJ/kmol)
1	$C + O_2 \rightarrow CO_2$	-394.9
2	$2C + O_2 \rightarrow 2CO$	-226.0
3	$2CO + O_2 \rightarrow 2CO_2$	-563.8
4	$C + 2H_2 \rightarrow CH_4$	-87.5
5	$C + H_2O \rightarrow CO + H_2$	+135.7
6	$C + CO_2 \rightarrow 2CO$	+168.9
7	$CO + H_2O \rightarrow CO_2 + H_2$	-33.2
8	$CO + 3H_2 \rightarrow CH_4 + H_2O$	-206.2
9	$CO_2 + 4H_2 \rightarrow CH_4 + 2H_2O$	-165.0

However, although carbon is the dominant species present in coal, the reactivity of coal may differ from that of pure carbon [7]. Hence, the coal gasification process is complicated by a number of factors, such as operational pressure, reaction temperature, coal rank and its composition and structure, and type of oxidants used [7, 15-18]. For instance, bituminous and anthracite coals usually have low moisture and volatile contents, hence less reactants available from within the coal available for gasification [19]. This was confirmed by Stańczyk et al. [20] demonstrating that gasification of lignite and hard coal blocks with air can lead to poor thermodynamic conditions and low process efficiency. Therefore, it is recommended to use oxygen and steam as main gasification agents to achieve high reaction temperatures and conversion of coal into syngas [19].

UCG experiments have been conducted worldwide for almost a century, both at in-situ and laboratory scale. The former ones demonstrated the feasibility of the UCG process and challenges that need to be overcome for successful commercialisation, such as process control, groundwater pollution and subsidence [21-28]. The later ones gave further confidence on the effectiveness and stability of the UCG process [8, 13, 14, 20, 29-38]. For instance, Stańczyk et al. [8] proved the feasibility of producing a H<sub>2</sub>-rich gas during the gasification of hard coal by alternating the injection of oxygen and steam. Liu et al. [37] showed that oxygen enriched air gasification of lignite is superior compared with air gasification producing a syngas with 2.4 higher calorific value. Similarly, higher flow rates of oxygen during the gasification of low rank coals led to higher gasification efficiency by increasing the reaction temperature and expansion of gasification area [35]. Overall, most laboratory studies have concentrated on the

influence of different gasification agents on the gasification of low and medium rank coals under atmospheric pressure conditions, which is potentially attributable to the complexity of gasification under high pressure regime but also limited reactivity and permeability of the high rank coals. This paper therefore aims to provide further understanding of the effects of various operating conditions on the CH<sub>4</sub> oriented coal gasification in the context of UCG by utilising two coals of different rank, i.e. bituminous coal and semi-anthracite. The experimental plan involved testing the influence of gaseous reactants (O<sub>2</sub> and steam), their supply rates (molar ratios of H<sub>2</sub>O:O<sub>2</sub>=1:1 and H<sub>2</sub>O:O<sub>2</sub>=2:1) and thermodynamic conditions (temperatures of 650°C, 750°C, 850°C and pressures of 20 bar and 36 bar) on the gasification process. In total, 20 gasification tests were conducted using a small-scale ex-situ gasification reactor. Based on a detailed analysis of experimental data obtained, the most favourable gasification conditions for CH<sub>4</sub>-rich gas production are identified. According to the authors' knowledge, no CH<sub>4</sub> focused gasification experiments in the context of UCG have been performed on coals of such rank. Hence, the dataset presented here offers insights into the most favourable thermodynamic conditions to maximise the CH<sub>4</sub> generation during gasification of bituminous and semi-anthracitic coals.

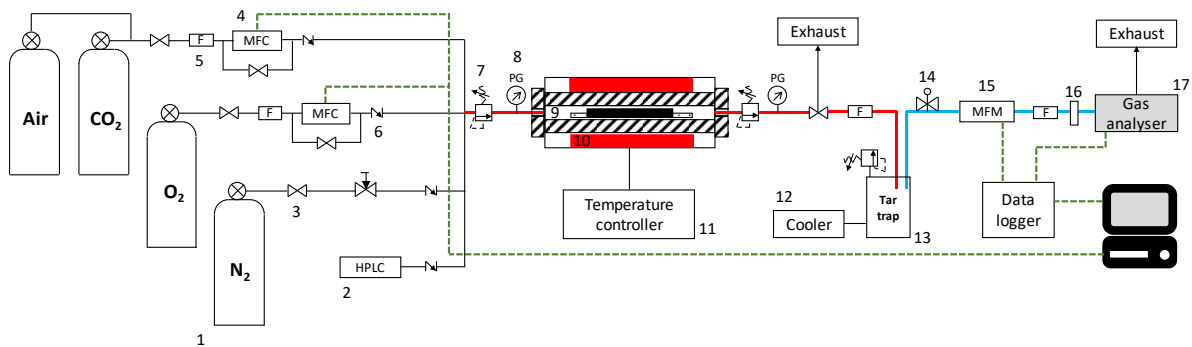
## 2. Materials and Methods

### 2.1. Experimental facility

The experimental simulations of the gasification process involved the use of a bespoke small-scale ex-situ high-pressure high-temperature gasification rig (Fig. 1). The rig can sustain maximum pressures and temperatures up to 50 bar and 900°C, respectively. Air, O<sub>2</sub>, CO<sub>2</sub> and N<sub>2</sub> can be used as reactant gases under supply rates up to 10 L/min and H<sub>2</sub>O under supply rates up to 9.99 mL/min.

The experimental setup consists of: 1) Gas cylinders with pressure regulators; 2) High Performance Liquid Chromatography (HPLC) water pump which supplies de-ionised water in the system; 3) On/off valves; 4) Mass flow controllers (MFC) with bypass on/off valve; 5) Filters; 6) Non-return valves; 7) Pressure relief valves; 8) Pressure gauges; 9) Tubular reactor 33 cm in length, 2.54 cm internal diameter; 10) Horizontal split hinge furnace; 11) Temperature controller Carbolite 301; 12) Water cooler

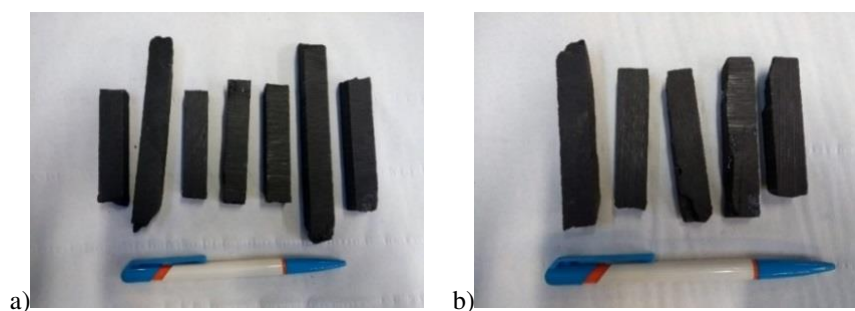
condenser; 13) Gas liquid separator - tar trap; 14) Back-pressure regulator; 15) Digital mass flow meter (MFM); 16) Rotameter and 17) Gas analyser.



**Fig. 1.** Schematic of the bespoke high-pressure high-temperature ex-situ small-scale gasification rig.

## 2.2. Materials

Field specimens were gathered from two different locations. The first selection of coal blocks was obtained from an open cast coal mine in the South Wales Coalfield, UK. The sampling location is 88 m below the ground level, from a coal seam of an average thickness of 1.2 m. The seam name is “Six Feet”, with a local name “Big”. The second selection of blocks was obtained from an underground coal mine Wesola in Poland. The sampling location is 950 m below the ground level from a coal seam of an average thickness of 5 m. Coal samples were cut out of the coal blocks using a saw with a diamond tipped saw blade. For the purpose of the gasification experiments, rectangular coal samples were cut to a length of 5-10 cm and a width of 1.5 cm (Fig. 2). Total mass of samples used in each experiment was approximately 30 g (Table 2). Crushed samples passed through a sieve size of 0.212 mm were used for the Proximate and Ultimate analyses [39-43], and the results are presented in Table 3. An air-drying method was applied to dry the samples prior to the experiments [44].



**Fig. 2.** Coal samples used in the gasification tests: a) “Six Feet” coal; b) “Wesola” coal.

129 **Table 2.** Mass of coal samples used for ex-situ small-scale gasification tests.

Test number	Mass (g)	
	“Six Feet” coal (South Wales, UK)	“Wesoła” coal (Silesia, Poland)
Test 1	30.96	30.52
Test 2	30.31	30.67
Test 3	31.35	30.89
Test 4	30.86	30.95
Test 5	30.91	30.82
Test 6	30.78	30.81
Test 7	30.25	30.21
Test 8	30.21	30.55
Test 9	30.56	30.70
Test 10	30.88	30.94
Average	30.71	30.71

130 **Table 3.** Proximate and Ultimate characteristics of coals used for ex-situ small-scale gasification tests.

Parameter	Value (%)	
	“Six Feet” coal (South Wales, UK)	“Wesoła” coal (Silesia, Poland)
Moisture content	0.88	3.60
Ash content	2.99	8.74
Volatile matter	13.42	27.67
Total carbon content	89.06	75.35
Total sulphur content	0.70	0.31

### 131 2.3.Experimental methodology

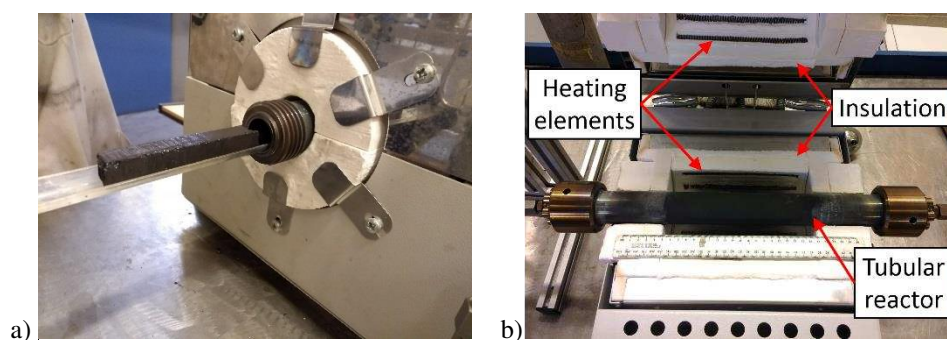
132 Ten tests on each coal were conducted using mixtures of O<sub>2</sub> and H<sub>2</sub>O as the gasification media under  
133 different molar ratios, pressure and temperature conditions. In particular, molar ratios of H<sub>2</sub>O:O<sub>2</sub>=1:1  
134 and H<sub>2</sub>O:O<sub>2</sub>=2:1, pressures of 20 bar and 36 bar and temperatures of 650°C, 750°C and 850°C were  
135 considered in the gasification tests (Table 4).

136

137 **Table 4.** Experimental conditions used for ex-situ small-scale gasification tests.

Test No.	Temperature (°C)	Pressure (bar)	O <sub>2</sub> flow rate (L/min)	H <sub>2</sub> O flow rate (mL/min)	H <sub>2</sub> O:O <sub>2</sub> molar ratio
Test 1	650	20	0.4	0.3	1:1
Test 2	650	20	0.4	0.6	2:1
Test 3	850	20	0.4	0.3	1:1
Test 4	850	20	0.4	0.6	2:1
Test 5	650	36	0.7	0.53	1:1
Test 6	650	36	0.7	1.05	2:1
Test 7	850	36	0.7	0.53	1:1
Test 8	850	36	0.7	1.05	2:1
Test 9	750	20	0.4	0.6	2:1
Test 10	750	36	0.7	1.05	2:1

138 Prior to each experiment, a sample was loaded in a quartz boat which was inserted into the tubular  
 139 pressure reactor (Fig. 3). Upon closing the pressure reactor and connecting it to the gas pipeline, the  
 140 system was isolated by closing the valves. The pressure of the system was then raised up to the  
 141 designated value using O<sub>2</sub> by applying a flow rate of 3 l/min. Once the pressure stabilised, the flow rate  
 142 was reduced to the predetermined value (see Table 4) and the valves were opened. The temperature of  
 143 the system was then increased up to the predetermined value setting up a heating rate of 20 °C/min.  
 144 Injection of de-ionised water at a predetermined flow rate (see Table 4), via an HPLC pump,  
 145 commenced upon temperature reaching 200°C which was then evaporated to steam due to the high  
 146 temperature in the system ensured by the temperature controller providing a constant heat to the reactor.  
 147 Once the temperature reached the predetermined value, each experiment was run at a constant  
 148 temperature for 90 minutes.

149 **Fig. 3.** Gasification rig assembly: a) Sample insertion; b) Tubular reactor with heating elements and insulation.



150 Hence, the gasification process was allothermal where the heat required for gasification reactions was  
151 produced outside of the reactor, via temperature controller, and then transferred inside. This allowed  
152 for an assessment of coal gasification at a range of temperatures which were controlled in a precise  
153 manner, opposite to the autothermal gasification which lacks an inside temperature control.

154 Eight tests (Tests 1-8) with experimental conditions described in Table 4 were conducted first. Based  
155 on the observations, which will be explained in the following section, two more tests (Test 9 and Test  
156 10) were conducted to further analyse the impact of temperature on gas concentration with a purpose to  
157 obtain the most favourable conditions for producing a CH<sub>4</sub>-rich syngas.

### 3. Results and discussion

This section presents the results of gasification experiments conducted both on the “Six Feet” coal and “Wesoła” coals. It is noted that the figures presented in the following sections contain absolute average gas concentration values in vol.%, serving as a basis for the analysis in which positive or negative changes of gas concentrations as a result of changes in the pressure, temperature, reactant supply rates and coal rank are discussed.

#### 3.1. Product gas composition

The time evolutions of product gas compositions during gasification experiments conducted on the “Six Feet” and “Wesoła” coals are presented in Fig. 4 and Fig. 5, respectively. For experiments where the temperature was raised to 650°C, maximum CH<sub>4</sub> concentrations can be observed in the period when the temperature had already reached 650°C. For experiments to be undertaken at 850°C, such peaks in CH<sub>4</sub> concentrations are obtained during the temperature ramping period, i.e. in the zone below 850°C. In particular, the CH<sub>4</sub> concentration peaks were achieved and prolonged for a certain period within the temperature zone of 650-780°C and 560-800°C for the “Six Feet” and “Wesoła” coals, respectively. Based on those observations, it was decided that two more tests, i.e. Test 9 and Test 10, are performed under the temperature condition of 750°C (using 20 bar and 36 bar pressures at H<sub>2</sub>O:O<sub>2</sub>=2:1 molar ratio) in order to provide a verification of the results and confirm the optimal temperature to obtain a steady production of CH<sub>4</sub>-rich syngas.

The results obtained on “Six Feet” coal reveal that CH<sub>4</sub> concentration started to rapidly increase after approximately 20-25 minutes from the start of each test conducted at 20 bar when the temperature reached between 350-460°C (Fig. 3). For the “Wesoła” coal, it took 18-22 minutes for the CH<sub>4</sub> production to occur when the temperature reached 300-460°C, at the same pressure conditions (Fig. 4). In general, CH<sub>4</sub> formation begins between 300°C and 550°C and as the temperature continues to rise, semi-coke begins to solidify and contract followed with the yield of CH<sub>4</sub>, H<sub>2</sub> and CO<sub>2</sub> [2, 17]. The fact that CH<sub>4</sub> formation starts earlier for the “Wesoła” coal compared to “Six Feet” coal might be related to coal pyrolysis which is influenced by the coal rank. Liu et al. [17] have shown that temperatures at

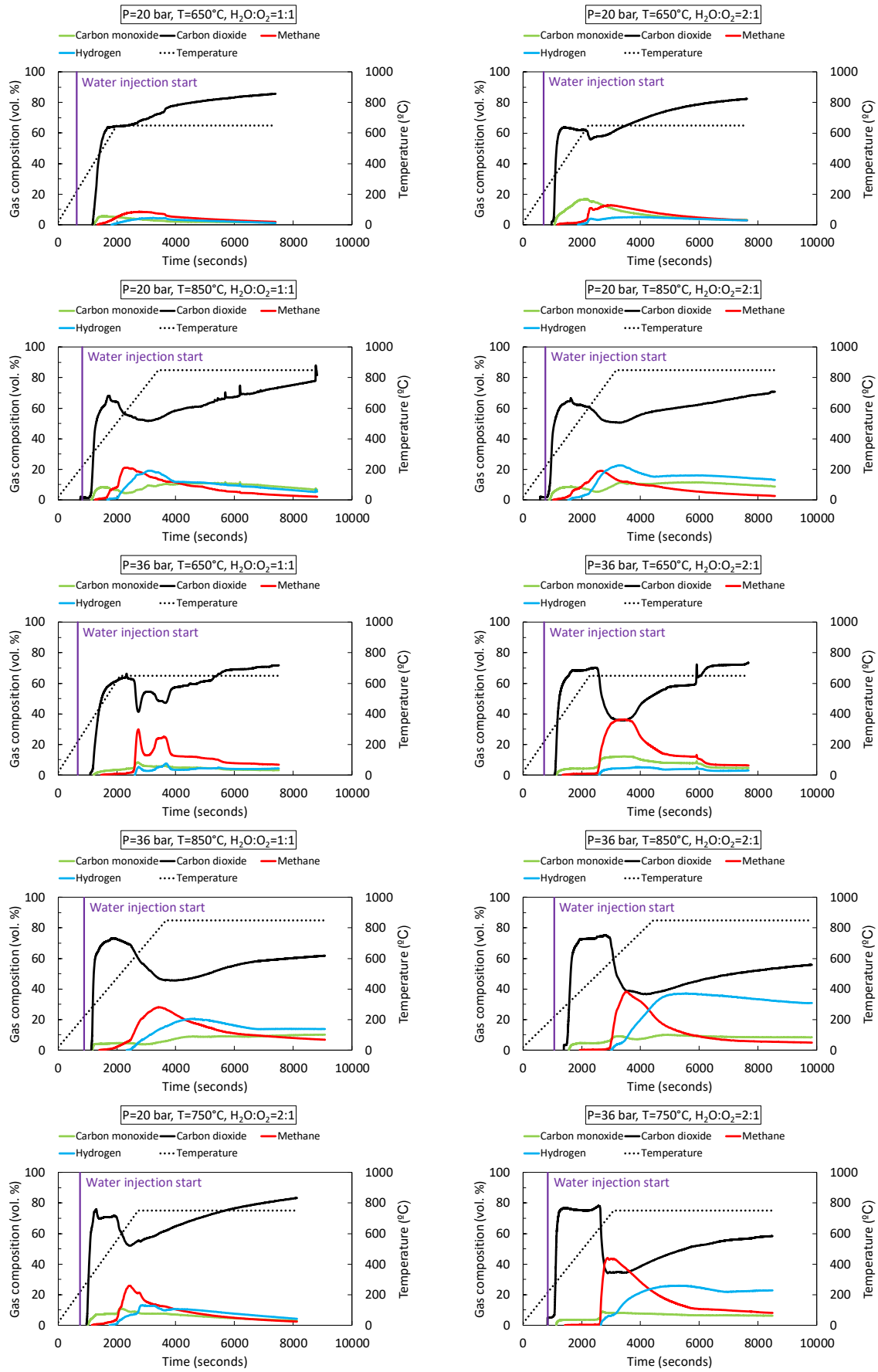
which CH<sub>4</sub> forms shift towards high temperatures with an increase in coal rank, as more active groups and radicals are available in lower rank coals increasing the instantaneous formation rate of CH<sub>4</sub> inside the coal particles.

It can be inferred that the increase in pressure from 20 bar to 36 bar as well as the increase in the amount of steam with respect to oxygen increase the time required for the beginning of CH<sub>4</sub> production. This is reasonable to expect as changes in the pressure can shift the chemical equilibrium of the system as well as affect the heat and mass transfer, while excess steam in the system can affect the thermal efficiency of the process [7, 15].

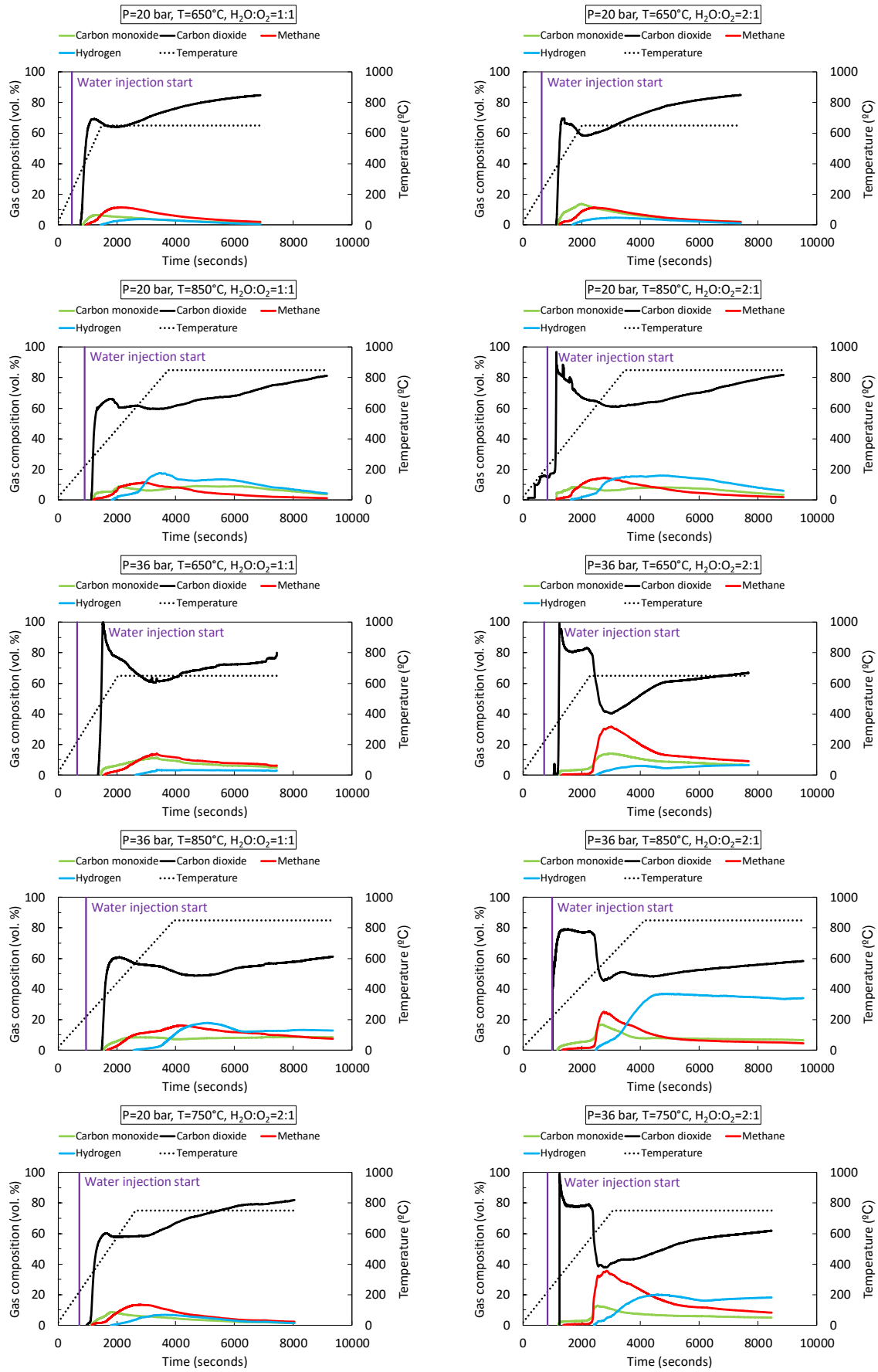
CO occurs earlier than H<sub>2</sub> both at experiments conducted at 20 bar and 36 bar, i.e. 10-16 minutes and 20-26 minutes earlier for the “Six Feet” coal and 6-12 minutes and 16-22 minutes earlier for the “Wesoła” coal, respectively. Early breakthrough of CO is due to the partial oxidation of coal, followed by endothermic reactions favoured by an increase in temperature, i.e. Boudouard reaction and steam gasification producing more CO and H<sub>2</sub>. Large volumes of H<sub>2</sub> will only be produced at temperatures much higher than 600°C as the high temperature is required for the C-H bonds to rupture [17, 45].

CO<sub>2</sub> is the first gas produced in each test, owing to strong oxidation of coal, which shows a constant increase throughout each experiment, i.e. initial rapid increase in concentration, followed by a drop in CO<sub>2</sub> concentration which coincides with peak production of CH<sub>4</sub>, H<sub>2</sub> and CO after which gradual increase of CO<sub>2</sub> until the end of each experiment occurs. This is related to the fact that oxidising reactions continue as long as there is free O<sub>2</sub> in the injection stream, yielding high amounts of CO and CO<sub>2</sub> [16].

A summary of average gas compositions obtained for the duration of each experiment conducted on both coals is given in Table 5. Based on the results for “Six Feet” coal, the highest average CH<sub>4</sub> concentration of 15.34 vol.% was observed in Test 10 (T=750°C, P=36 bar, H<sub>2</sub>O:O<sub>2</sub>=2:1). Under the same experimental conditions, the highest average CH<sub>4</sub> concentration of 14.64 vol.% was obtained during the gasification of “Wesoła” coal. The peak CH<sub>4</sub> concentrations observed for the “Six Feet” and “Wesoła” coals were 44.00 vol.% and 35.55 vol.%, respectively.



**Fig. 4.** Changes in gas compositions obtained during the experiments conducted on “Six Feet” coal samples at different experimental conditions.



**Fig. 5.** Changes in gas compositions obtained during the experiments conducted on "Wesola" coal samples at different experimental conditions.

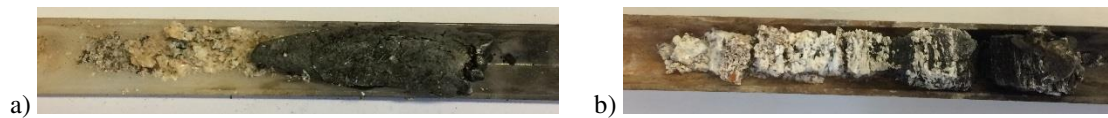
214 **Table 5.** Average gas concentrations and calorific values obtained in tests conducted on both coals.

Test No	T (°C)	P (bar)	H <sub>2</sub> O:O <sub>2</sub> ratio	„Six Feet” coal					„Wesola” coal				
				Average gas concentration (vol. %)				Calorific value (MJ/Nm <sup>3</sup> )	Average gas concentration (vol. %)				Calorific value (MJ/Nm <sup>3</sup> )
				CH <sub>4</sub>	H <sub>2</sub>	CO	CO <sub>2</sub>		CH <sub>4</sub>	H <sub>2</sub>	CO	CO <sub>2</sub>	
1	650	20	1:1	4.45	2.61	2.63	74.14	2.21	5.88	2.23	3.30	73.24	2.77
2	650	20	2:1	6.36	3.86	7.40	69.16	3.63	5.52	2.94	5.99	72.22	3.06
3	850	20	1:1	7.89	9.87	8.80	60.92	5.01	4.81	9.90	7.01	67.33	3.68
4	850	20	2:1	7.36	14.56	9.54	57.83	5.42	6.53	11.06	6.61	63.97	4.37
5	650	36	1:1	9.85	4.10	4.20	59.26	4.51	8.30	2.84	7.34	69.63	4.21
6	650	36	2:1	13.70	3.74	7.20	57.94	6.23	13.56	5.16	8.30	60.60	6.47
7	850	36	1:1	12.48	14.47	7.56	56.78	7.00	10.70	11.64	7.81	54.52	6.08
8	850	36	2:1	11.16	29.87	7.99	50.29	8.24	8.10	30.14	7.78	56.58	7.14
9	750	20	2:1	7.89	7.83	5.92	69.33	4.42	6.39	3.80	3.93	67.12	3.20
10	750	36	2:1	15.34	20.94	6.31	52.01	8.56	14.64	15.65	6.27	56.32	7.73

215

However, Test 10 produced less  $H_2$ , and CO than Test 8 during the gasification of both coals. This proves that the most favourable conditions for the methanation reaction are at 750 °C, in which hydrogen and carbon monoxide are consumed to form methane. Owing to high methane content obtained in Test 10, average calorific values of the syngas produced in Test 10 were 4% and 8% higher than in Test 8 for the “Six Feet” and “Wesoła” coals, respectively.

Fig. 6 and Table 6 show the gasification residues and their masses, respectively. It can be observed that the gasification proceeds in one direction and the ash layer (white area) forms at the end where reactants are injected. By comparing the results presented in Table 6 and Table 2, about 2/3 of the total coal mass for both coals were consumed by chemical reactions. However, one can observe that 12% less of gasification residue were left after the gasification of “Wesoła” coal compared to “Six Feet” coal.



**Fig. 6.** Gasification residues: a) “Six Feet” coal; b) “Wesoła” coal.

**Table 6.** Mass of the gasification solid residues obtained from ex-situ small-scale gasification tests.

Test number	Mass (g)	
	“Six Feet” coal	“Wesoła” coal
Test 1	12.51	9.02
Test 2	11.13	9.33
Test 3	8.18	4.29
Test 4	7.92	5.35
Test 5	16.52	14.60
Test 6	8.16	16.78
Test 7	13.21	14.28
Test 8	13.20	11.68
Test 9	11.51	6.64
Test 10	18.18	13.63
Average	12.05	10.56

### 3.2.Effect of pressure

The effect of pressure on the reaction rate depends on the specific reaction involved [7]. Some gasification reactions in which the number of moles of reactants is higher than the number of moles of products, e.g. hydrogasification and methanation reactions, are favoured at high pressures [12]. Increasing the pressure of the system increases the average  $CH_4$  concentrations in all experiments. For instance, in experiments conducted on “Six Feet” coal at 650°C, 750°C and 850°C with  $H_2O:O_2=2:1$ , a

change in pressure from 20 to 36 bar increases the CH<sub>4</sub> concentration by 115% and 94%, 52% respectively. Similarly, raising the pressure yields 146%, 129% and 24% increase in CH<sub>4</sub> concentration during gasification of “Wesoła” coal at 650°C, 750°C and 750°C with H<sub>2</sub>O:O<sub>2</sub>=2:1, respectively.

Although it is generally suggested in the literature that the production of CO<sub>2</sub> is favoured at higher pressures [12], this is not observed in this work which could be due to the complexity of the principal chemical reactions between carbon and oxygen that occur simultaneously and sequentially. As Lee et al. [7] suggested, oxidation of coal is significantly more complex than oxidation of pure carbon and the mechanistic chemistry explained purely through the equations in Table 1 may not be sufficient, as coal contains various reactive organic functional groups and some mineral ingredients which act as catalysts. One of the reasons may also be the intensification of the CO<sub>2</sub> methanation under the conditions of experiments conducted. During the gasification of “Six Feet” and in particular the “Wesoła” coal, a general trend is visible where increase in pressure generally has a positive impact on CO, especially at 650°C and H<sub>2</sub>O:O<sub>2</sub>=1:1, increasing the concentrations by 60% and 122%, respectively. This may be related to the fact that Boudouard reaction requires high pressure for significant conversion for higher reactant concentrations [7].

### 3.3.Effect of temperature

Increase in temperature above 650°C has a positive effect on CH<sub>4</sub> generation. In particular, as the temperature increases from 650°C to 850°C, CH<sub>4</sub> concentration increases by up to 77% for the “Six Feet” coal (at H<sub>2</sub>O:O<sub>2</sub>=1:1 and 20 bar) and 29% for the “Wesoła” coal (at H<sub>2</sub>O:O<sub>2</sub>=1:1 and 36 bar). However, for both coals under study, syngas with the highest CH<sub>4</sub> concentration was obtained at 750°C. Hydrogasification, methanation and volatile matter are the main sources for CH<sub>4</sub> generation [36]. Furthermore, hydrogasification and methanation reactions are exothermic and favoured at temperatures ≤670°C, opposite to steam and CO<sub>2</sub> gasification reactions [7]. Hence, as the temperature increases, they will proceed at lower rate decreasing the CH<sub>4</sub> production [7]. Increase in CH<sub>4</sub> concentration with temperature could also be related to the pyrolysis of coal during which CH<sub>4</sub> is mainly released within the temperature zone of 450-600°C, predominantly by the breakage of methylene bridges [46, 47].



However, although CH<sub>4</sub> can be formed at temperatures  $\geq 700^{\circ}\text{C}$  due to secondary devolatilisation, only small amounts of CH<sub>4</sub> are produced at such temperatures as a result of pyrolysis [17]. Increase in CH<sub>4</sub> concentration at temperatures higher than 700°C is predominantly due to excess H<sub>2</sub> in the system that participates in the hydrogasification reaction, as high hydrogen pressure significantly enhances this reaction [7, 17].

Both CO and H<sub>2</sub> concentrations show a strong correlation with temperature, increasing up to 235% and 699% for “Six Feet” coal and up to 112% and 484% for the “Wesoła” coal, respectively. High temperature ( $>680^{\circ}\text{C}$ ) benefits the Boudouard reaction enhancing the CO production, similar to the steam gasification reaction [7, 36]. The water gas shift reaction also contributes to the generation of H<sub>2</sub> and CO<sub>2</sub>, but it is least dependent on temperature [7]. During pyrolysis, H<sub>2</sub> evolution starts at 400°C due to the condensation of aromatic structures and it progresses at a relatively high rate up to 1000°C, contributing continuously to the overall H<sub>2</sub> content in the syngas [45-47]. It is commonly understood that O<sub>2</sub> reacts completely in a very short distance from the location at which it comes in contact with coal, which is further pronounced at higher temperatures as the rates of reactions are intrinsically higher [7, 12]. Hence, as the reaction pathway is very complex involving several carbon-oxygen interactions, there is a general agreement that the ratio of CO to CO<sub>2</sub> increases with temperature [7, 45], as also supported through this work.

### 3.4. Effect of gasification reactant supply rates

Generation of CH<sub>4</sub> is generally favoured by doubling the number of moles of steam with respect to O<sub>2</sub> in the system, which is particularly visible at 650°C. For instance, at 36 bar during the gasification of “Six Feet” and “Wesoła” coals the CH<sub>4</sub> concentration increased by 39% and 63%, respectively.

Similar can be observed for CO and H<sub>2</sub>, as it increased by up to 181% and 106% during “Six Feet” coal gasification and up to 82% and 159% during “Wesoła” coal gasification, respectively.

The observed behaviour is driven by the steam gasification and water gas shift reactions which are favoured by an excess amount of steam. The former one is an endothermic reaction requiring heat input, while the latter one is mildly exothermic and takes place on the solid surface of the coal where it is

being further catalysed by carbon surfaces, despite all four chemical species involved in the reaction are in gaseous state [7]. Produced  $H_2$  then primarily participates in the hydrogasification reaction resulting in an increased production of  $CH_4$ , but can also play role in the Sabatier reaction in which it reacts with  $CO_2$  to generate  $CH_4$  and steam [36]. Hence, high yield of  $CH_4$  is controlled by the molar ratio of  $H_2$  to CO or  $CO_2$ , depending on the reaction, suggesting that adjusting the  $H_2$  content in the syngas through steam injection and temperature control can benefit  $CH_4$  production.

### 3.5. Effect of coal rank

The gasification of coals of different rank yields notable differences in average concentrations of various gas species. The production of  $CH_4$  was consistently higher during the gasification of the “Six Feet” semi-anthracitic coal compared to the “Wesola” bituminous coal, showing up to 64% higher concentration, depending on the experimental conditions applied. Similarly,  $H_2$  and CO concentrations were generally higher during the gasification of the “Six Feet” than “Wesola” coal, i.e. up to 50% and 106% higher, respectively. Although the effect of coal rank does not have a pronounced effect on average concentrations of  $CO_2$  produced, the values obtained suggest a mild tendency to decrease with an increase in coal rank.

In general, the gasification of coal is affected by its properties such as carbon content, volatile matter, ash content, pore structure, particle size and thermodynamic conditions [7, 48]. Coals of higher rank can generate more  $CH_4$ , especially during the pyrolysis process, as the loss of oxygen-functional sites during the maturation process can increase the concentration of the  $CH_4$  precursors [17]. Furthermore, high effective carbon content during the stable gas production phase enforces the reduction of  $CO_2$  and increases the CO content [16]. Coals of higher rank contain predominantly micropores which offer most of the surface area where the actual gasification takes place in comparison to lower rank coals which consist of macropores, mesopores and micropores [18, 49]. However, as meso- and macropores offer passage to gasification agent to reach the reactant sites in the micropores, lower rank coals are more reactive [18].

## 4. Conclusions

The influence of gaseous reactants, i.e.  $O_2$  and steam, and their supply rates at different temperatures and pressures on the gasification process of “Six Feet” semi-anthracitic coal from the South Wales coalfield (UK) and “Wesoła” bituminous coal from the Silesian basin (Poland) to find the favourable experimental conditions for  $CH_4$ -rich syngas production was presented in this paper. The following conclusions can be made:

- Experiments where the temperature of the system was increased to  $750^\circ C$  demonstrated that it is the most favourable temperature around which the highest peak was achieved and sustained for a certain period of time compared to experiments conducted at  $650^\circ C$  and  $850^\circ C$ .
- Increasing the pressure from 20 bar to 36 bar increased the  $CH_4$  concentration. This was the most pronounced in experiments conducted at  $750^\circ C$  and  $H_2O:O_2=2:1$ , where  $CH_4$  concentration increased by 94% and 129% for “Six Feet” coal and “Wesoła” coal, respectively.
- Doubling the number of moles of steam with respect to  $O_2$  in the system generally increased the  $CH_4$  concentration, by up to 43% and 63% for “Six Feet” coal and “Wesoła” coal, respectively.
- By increasing the temperature of the system from  $650^\circ C$  to  $850^\circ C$ ,  $CH_4$  concentration increased by up to 77% and 29% for “Six Feet” Coal and “Wesoła” coal, respectively. However, through experiments on both coals, it was demonstrated that the optimal temperature for stable and  $CH_4$ -rich syngas production is  $750^\circ C$  at 36 bar and  $H_2O:O_2=2:1$ .
- The highest peak and average  $CH_4$  concentrations of 44.00 vol.% and 15.34 vol.% were observed during the gasification of “Six Feet” coal, and concentrations of 35.55 vol.% and 14.64 vol.% during the gasification of “Wesoła” coal, respectively. This demonstrated that an increase in coal rank favours the  $CH_4$  generation.
- Overall, as the  $CH_4$  production was favoured by an increase in pressure and an increase in the amount of steam in the system, and keeping the temperature at  $750^\circ C$  generated syngas with the highest average  $CH_4$  content, the amount of steam injected and the dynamics of steam and

O<sub>2</sub> supply rates at high pressures can be the controlling factors through which the syngas with high CH<sub>4</sub> content is produced.

- Based on the findings of this work, generation of CH<sub>4</sub>-rich syngas during in situ gasification could be achieved by selecting deep carbon-rich coal seams to enable gasification at high pressure conditions. In order to maintain the gasification at temperatures favourable for CH<sub>4</sub> production which is directly dependent on the amount of water and hydrogen amounts available within the reactor, hydrogeology of the site would need to be carefully assessed and understood so that the groundwater flowing towards the reactor would be provided at a sufficient rate to adsorb the heat within the reactor and converts to steam. This work suggests that doubling the amount of water with respect to the primary gasification agent (oxygen or air) would be beneficial and serve as a starting point, however, the optimal amount would need to be assessed and adjusted based on the quality of the product gas.

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